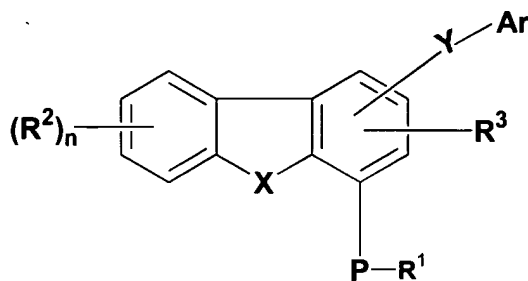


LISTING OF THE CLAIMS

This listing of claims will replace all prior versions, and listings, of the claims in the application.

1. (currently amended) A compound of general formula (1)



(1)

wherein:

R^1 , R^2 and R^3 may be same or different and are independently selected from the groups consisting of hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heterocyclic ring, substituted or unsubstituted heterocyclalkyl, substituted or unsubstituted heteroarylalkyl, $-C(O)-R^1$, $-C(O)O-R^1$, $-C(O)NR^1R^1$, $-S(O)_m-R^1$, $-S(O)_m-NR^1R^1$, nitro, $-OH$, cyano, amino, formyl, acetyl, halogen, $-OR^1$, halogen, $-OR^1$, $-SR^1$, or a protecting groups group ~~or and~~ when two R^2 substituents are ortho to each other, they may be joined to $[[a]]$ form a saturated or unsaturated cyclic ring, which may optionally include up to two heteroatoms selected from O, NR^1 or S;

~~wherein~~ P represents is oxygen or sulfur;

~~wherein~~ n represents is an integer from 0 – 4;

alkylcarbonyloxy' ~~and~~ may be the same or different ~~which and are~~ one or more of selected from the ~~groups such as~~ hydrogen, hydroxy, halogen, carboxyl, cyano, nitro, oxo (=O), ~~thio thio~~(=S), substituted or unsubstituted alkyl, substituted or unsubstituted alkoxy, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted amino, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, [[']]substituted heterocyclalkyl ring[[']] , substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted heterocyclic ring, substituted or unsubstituted guanidine, -COOR^x, -C(O)R^x, -C(S)R^x, -C(O)NR^xR^y, -C(O)ONR^xR^y, -NR^xCONR^yR^z, -N(R^x)SOR^y, -N(R^x)SO₂R^y, ~~(=N-N(R^x*)R^y)~~ =N-N(R^x)(R^y), -NR^xC(O)OR^y, -NR^xC(O)OR^y; -NR^xR^y, -NR^xC(O)R^y-, -NR^xC(S)R^y-, -NR^xC(S)NR^yR^z-, -SONR^xR^y-, -SO₂NR^xR^y-, -OR^x, -OR^xC(O)NR^yR^z-, -OR^xC(O)OR^y-, -OC(O)R^x, -OC(O)NR^xR^y, -R^xNR^yC(O)R^z, -R^xOR^y, -R^xC(O)OR^y, -R^xC(O)NR^yR^z, -R^xC(O)R^x, -R^xOC(O)R^y, -SR^x, -SOR^x, -SO₂R^x, or -ONO₂, wherein R^x, R^y and R^z ~~in each of the above groups can be~~ are independently hydrogen atom, substituted or unsubstituted alkyl, substituted or unsubstituted alkoxy, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted amino, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, [[']]substituted heterocyclalkyl ring[[']] substituted or unsubstituted heteroarylalkyl, substituted or an unsubstituted heterocyclic ring[[,]].

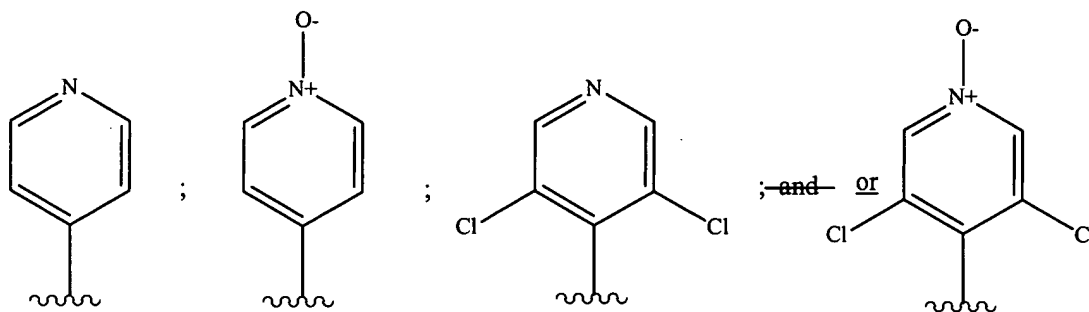
3. (original) The compound according to claim 1 wherein R¹ is substituted alkyl.
4. (original) The compound according to claim 3 wherein R¹ is CHF₂.
5. (original) The compound according to claim 1 wherein R¹ is unsubstituted alkyl.
6. (original) The compound according to claim 5 wherein R¹ is methyl.

7. (currently amended) The compound according to claim 1 ~~claims 1-5 or 6~~ wherein P is O or S.
8. (currently amended) The compound according to claim 7 ~~where~~ wherein P is O.
9. (currently amended) The compound according to claim 1 ~~claims 1-7 or 8~~ wherein R² is ~~selected from the group consisting of~~ substituted alkyl, halogen, cyano, nitro, amino, substituted heterocyclic ring or ~~and~~ SO₂NR¹R¹ and n=1.
10. (original) The compound according to claim 9 wherein R² is chloro.
11. (original) The compound according to claim 9 wherein R² is substituted alkyl.
12. (original) The compound according to claim 11 wherein R² is CF₃.
13. (original) The compound according to claim 9 wherein R² is -NH₂.
14. (currently amended) The compound according to claim 9 wherein R² is -SO₂NR¹R¹ - SO₂NR¹R².
15. (original) The compound according to claim 14 wherein R² is SO₂N(CH₃)₂.
16. (currently amended) The compound according to claim 1 ~~claims 1-14 or 15~~ wherein Y is -C(O)NH-.
17. (currently amended) The compound according to claim 1 ~~claims 1-15 or 16~~ wherein Ar is ~~selected from the group consisting of~~ substituted or unsubstituted 4-pyridyl; substituted or unsubstituted 4-pyridyl-N-oxide; substituted or unsubstituted 3-pyridyl ~~3-pyridyl~~, substituted or unsubstituted 3-pyridyl-N-oxide ~~3-pyridyl N-oxide~~; substituted or unsubstituted 2-pyridyl ~~2-pyridyl~~; ~~and or~~ substituted or unsubstituted 2-pyridyl N-oxide ~~2-pyridyl N-oxide~~.

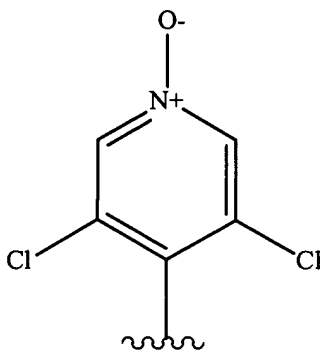
18. (currently amended) The compound according to claim 17 wherein said substituent Ar is substituted with halogen.

19. (original) The compound according to claim 18 wherein said halogen is chloro.

20. (currently amended) The compound according to claim 17 wherein Ar is selected from the group consisting of

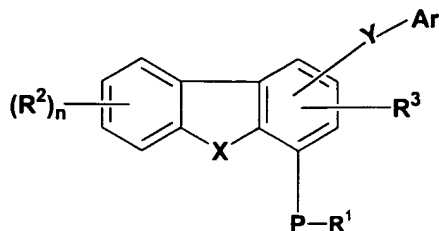


21. (original) The compound according to claim 20 wherein Ar is



Claims 22-59. (canceled)

60. (currently amended) A process for the preparation of a compound ~~compounds~~ of general formula (1)



(1)

wherein R¹, R² and R³ may be same or different and are independently selected from the groups consisting of hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heterocyclic group, substituted or unsubstituted heterocyclylalkyl, substituted or unsubstituted heteroarylalkyl, -C(O)-R¹, -C(O)O-R¹, -C(O)NR¹R¹, -S(O)_m-R¹, -S(O)_m-NR¹R¹, nitro, -OH, cyano, amino, formyl, acetyl, halogen, OR¹ halogen, -OR¹, -SR¹, or a protecting group groups ~~or~~ and when two R² substituents ortho to each other, they may be joined to a form a saturated or unsaturated cyclic ring, which may optionally include up to two heteroatoms selected from O, NR¹ or S;

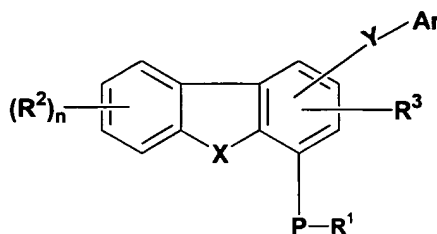
wherein P represents is oxygen or sulfur;

wherein n represents is an integer from 0 – 4;

Ar is substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, a substituted or unsubstituted heterocyclic ring or a substituted or unsubstituted heteroaryl ring;

X is oxygen, S(O)_m or NR⁵;

R⁵ is ~~represents~~ hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heterocyclic group, substituted or unsubstituted heterocyclalkyl, substituted or unsubstituted heteroarylalkyl, -C(O)-R¹, -C(O)O-R¹, -C(O)NR¹R¹, -S(O)_m-R¹, -S(O)_m-NR¹R¹, nitro, -OH, cyano, amino, formyl, acetyl, ~~halogen, -OR²~~ halogen, -OR², -SR² ~~and or~~ a protecting groups group;



(1)

wherein:

R¹, R² and R³ may be same or different and are independently selected from the groups consisting of hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heterocyclic group, substituted or unsubstituted heterocyclalkyl, substituted or unsubstituted heteroarylalkyl, -C(O)-R¹, -C(O)O-R¹, -C(O)NR¹R¹, -S(O)_m-R¹, -S(O)_m-NR¹R¹, nitro, -OH, cyano, amino, formyl, acetyl, halogen, OR¹ halogen, -SR¹, or a protecting group groups or and when two R² substituents are ortho to each other, they may be joined to form a saturated or unsaturated cyclic ring, which may optionally include up to two heteroatoms selected from O, NR¹ or S;

wherein P represents is oxygen or sulfur;

wherein n represents is an integer from 0 – 4;

Ar is substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, a substituted or unsubstituted heterocyclic ring or a substituted or unsubstituted heteroaryl ring;

X is oxygen, S(O)_m or NR⁵;

R⁵ represents is hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heterocyclic group, substituted or unsubstituted heterocyclalkyl, substituted or unsubstituted heteroarylalkyl, -C(O)-R¹, -C(O)O-R¹, -C(O)NR¹R¹, -S(O)_m-R¹, -S(O)_m-NR¹R¹, nitro,

-OH, cyano, amino, formyl, acetyl, ~~halogen, -OR²~~ halogen, -OR², -SR² ~~and or a~~ protecting groups
group;

m is 0, 1 or 2;

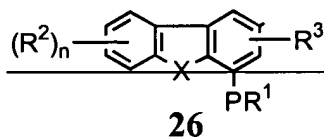
Y is -C(O)NR⁴, -NR⁴SO₂, -SO₂NR⁴ or -NR⁴C(O);

R⁴ is hydrogen, substituted or unsubstituted alkyl, hydroxyl, -OR¹, -COOR¹, substituted or unsubstituted aryl, or a substituted or unsubstituted heterocyclic ring ;

~~and their analogs, their tautomers, their regioisomers, their stereoisomers, their enantiomers, their diastereomers, their polymorphs, their pharmaceutically acceptable salts, their N oxides, their pharmaceutically acceptable solvates and their pharmaceutical compositions containing them or a pharmaceutical acceptable salts thereof;~~

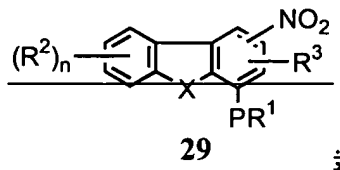
which comprises the steps of:

- (a) ~~nitrating the a~~ nitrating the a compound of general formula (26)

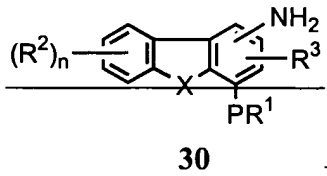


~~where the symbols are defined in the above~~

~~to yield the nitro compounds of general formula (29)~~



- (b) ~~reacting the compound of general formula (29) with a reducing agent to yield an the amino~~
~~compounds of general formula (30)~~



R^1 , R^2 and R^3 may be same or different and are independently ~~selected from the groups consisting of~~ hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl, substituted or unsubstituted unsubstituted alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heterocyclic group, substituted or unsubstituted heterocyclalkyl, substituted or unsubstituted heteroarylalkyl, $-C(O)-R^1$, $-C(O)O-R^1$, $-C(O)NR^1R^1$, $-S(O)_m-R^1$, $-S(O)_m-NR^1R^1$, nitro, $-OH$, cyano, amino, formyl, acetyl, halogen, $-OR^1$, $-SR^1$, ~~or a protecting group groups or~~ and when two R^2 substituents are ortho to each other, they may be joined to form a saturated or unsaturated cyclic ring, which may optionally include up to two heteroatoms selected from O, NR^1 or S;

wherein P represents is oxygen or sulfur;

wherein n represents is an integer from 0 – 4;

Ar is substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, a substituted or unsubstituted heterocyclic ring or asubstituted or unsubstituted heteroaryl ring;

X is oxygen, $S(O)_m$ or NR^5 ;

R^5 ~~represents~~ is hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl, substituted or ~~unsubstituted~~ unsubstituted alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted cycloalkylalkyl, substituted or unsubstituted cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted arylalkyl, substituted or unsubstituted heteroaryl, substituted or unsubstituted heterocyclic group, substituted or unsubstituted heterocyclalkyl, substituted or unsubstituted heteroarylalkyl, $-C(O)-R^1$, $-C(O)O-R^1$, $-C(O)NR^1R^1$, $-S(O)_m-R^1$, $-S(O)_m-NR^1R^1$, nitro, $-OH$, cyano, amino, formyl, acetyl, halogen, $-OR^2$, $-SR^2$ and or a protecting ~~groups group~~;

m is 0, 1 or 2;

Y is $-NR^4C(O)$;

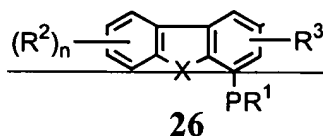
R^4 is hydrogen, substituted or unsubstituted alkyl, hydroxyl, $-OR^1$, $-COOR^1$, substituted or unsubstituted aryl, or a substituted or unsubstituted heterocyclic ring ;

~~and their analogs, their tautomers, their regioisomers, their stereoisomers, their enantiomers, their diastereomers, their polymorphs, their pharmaceutically acceptable salts, their N oxides, their~~

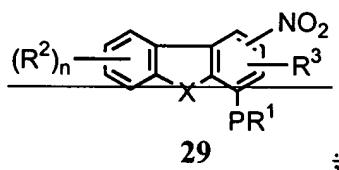
~~pharmaceutically acceptable solvates and their pharmaceutical compositions containing them or a pharmaceutical acceptable salts thereof;~~

which comprises the steps of;

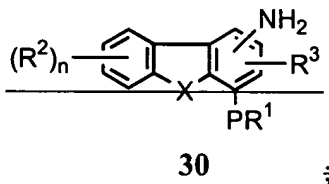
(a) ~~nitrating the compound of general formula (26)~~



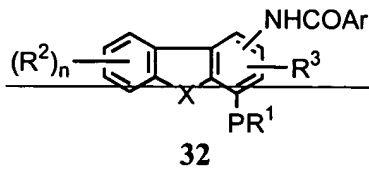
~~to yield the nitro compounds of general formula (29)~~



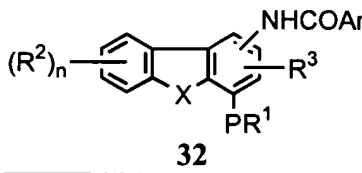
(b) ~~reacting the compound of general formula (29) with a reducing agent to yield the amino compounds of general formula (30)~~



(c) — reacting the amino compounds of general formula (30) with ArCOCl or a mixed anhydride of the formula ArCOOCOR^5 where R^5 -substituted or unsubstituted alkyl, cycloalkyl, aryl, heterocyclic ring, heteroaryl, to yield the compounds of general formula (32)



(d) alkylating a compound ~~the compounds~~ of general formula (32)



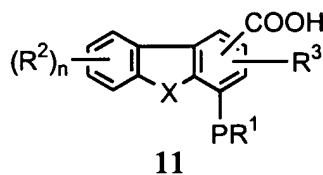
m is 0, 1 or 2;

Y is $-\text{C}(\text{O})\text{NR}^4$;

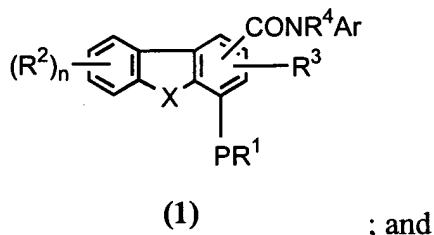
R^4 is hydrogen, substituted or unsubstituted alkyl, hydroxyl, $-\text{OR}^1$, $-\text{COOR}^1$, substituted or unsubstituted aryl, or a substituted or unsubstituted heterocyclic ring, or an N-oxide thereof;

comprising the steps of:

(a) reacting the compound of formula (11):



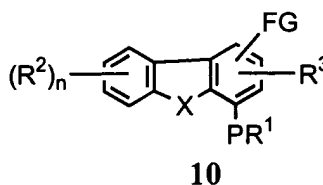
with an amine of the formula ArNHR^4 to yield a compound of formula (1)



(b) optionally converting the compound of formula (1) into its corresponding N-oxide.

75. (New) The method of claim 74 wherein the compound of formula (11) is formed by

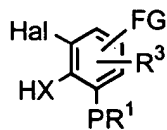
(a) converting the compound of general formula (10)



to general formula (11) wherein FG represents substituted or unsubstituted alkyl, formyl, cyano, halogen, nitro, amino or a carboxylic acid group.

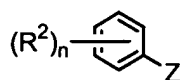
76. (New) The method of claim 75 wherein the compound of formula (10) is prepared by:

(i) reacting a compound of formula (13.a) with a compound of formula (23) under basic conditions



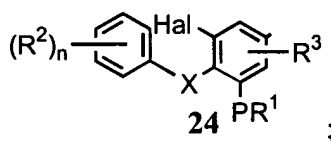
13.a

+



23

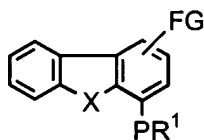
wherein Z is a halogen; FG is a substituted or unsubstituted alkyl, formyl, cyano, halogen, nitro, or amino; and Hal is halogen, to yield a compound of formula (24)



(ii) cyclizing the compound of general formula (24) under palladium catalyzed coupling conditions to form a tricyclic compound of general formula (10).

77. (New) The method of claim 75 wherein the compound of formula (10) is prepared by:

(i) reacting a compound of general formula (25) with an electrophile

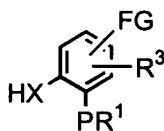


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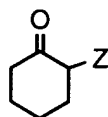
wherein FG is alkyl, formyl, cyano, halogen, nitro, or amino; to yield a compound of formula (10).

78. (New) The method of claim 75 wherein the compound of formula (10) is formed by:

(i) reacting a compound-of general-formula (13) with a compound of formula (20) under basic conditions

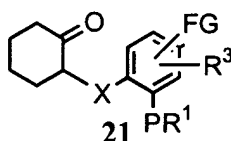


13



20

to yield a compound of general formula (21)

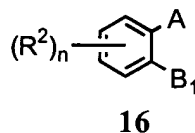


wherein FG is alkyl, formyl, cyano, halogen, nitro, or amino; and Z is a halogen; and

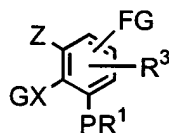
(ii) cyclizing the compound of general formula (21) under acidic conditions followed by oxidation to yield a tricyclic compound of general formula (10).

79. (New) The method of claim 75 wherein the compound of formula (10) is formed by:

(i) reacting a compound of formula (16) with a compound of formula (17)



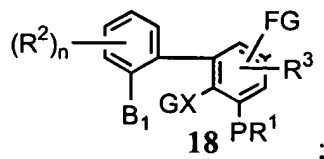
16



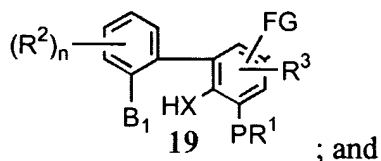
17

where A is halogen, -OMs, -OTs or -B(OH)₂; Ms is a methanesulfonyl group; Ts is a p-toluenesulfonyl group; B₁ is halogen; G is a protecting group selected from benzyloxycarbonyl, t-butyloxycarbonyl, isopropyl, cyclopentyl, allyl, acetyl and benzyl, FG is alkyl, formyl, cyano, halogen, nitro, or amino; and Z is halogen;

to yield a compound of formula (18)



- (ii) deprotecting the compound of formula (18) to yield a compound of formula (19)

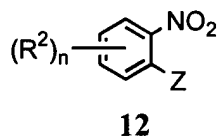


; and

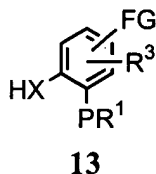
- (iii) cyclizing the intermediate of formula (19) under basic conditions to yield a tricyclic compound of formula (10).

80. (New) The method of claim 75 wherein the compound of formula (10) is prepared by:

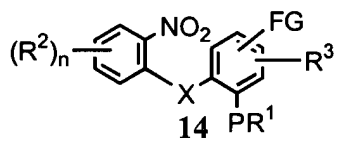
- (i) reacting a compound of general formula (12) where Z is a halogen



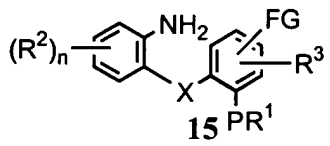
with an aromatic group of formula (13)



wherein FG is alkyl, formyl, cyano, halogen, nitro, or amino, under basic conditions to yield a compound of formula (14)



- (ii) reducing the compound of formula (14) to obtain a compound of formula (15)

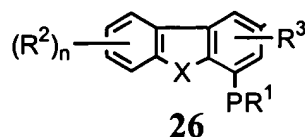


- (iii) cyclizing of the compound of formula (15) to yield a tricyclic compound of formula (10).

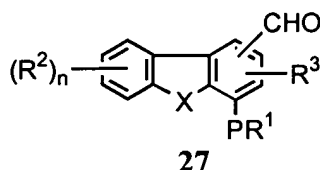
81. (New) The method of claim 75, wherein (i) FG is methyl and step (a) comprises oxidizing the compound of formula (10) with a manganese or chromium reagent to form a compound of

formula (11), (ii) FG is cyano and step (a) comprises hydrolyzing the compound of formula (10) to form a compound of formula (11), or (iii) FG is bromine and step (a) comprises reacting the compound of formula (10) with lithium followed by treatment with carbon dioxide to form a compound of formula (11).

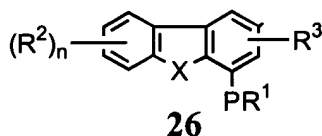
82. (New) The method of claim 74 wherein the compound of formula (11) is prepared by:
 (a) formylation of a compound of formula (26)



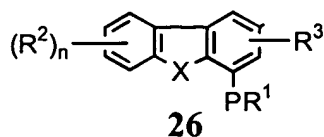
followed by oxidation of the aldehyde group in the resulting compound of formula (27)



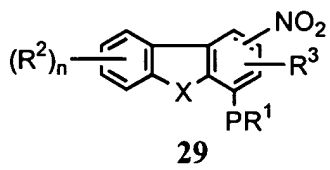
83. (New) The method of claim 60, wherein the compound of formula (28) is formed by chlorosulfonylation of the compound of formula (26)



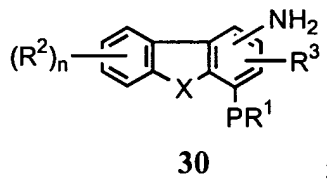
84. (New) The method of claim 61, wherein the compound of formula (31) is prepared by
 (a) nitrating a compound of formula (26)



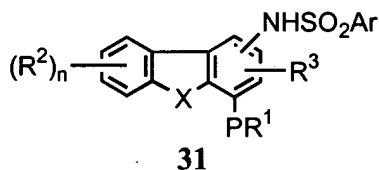
to yield a nitro compound of formula (29)



- (b) reacting the compound of formula (29) with a reducing agent to yield an amino compound of general formula (30)

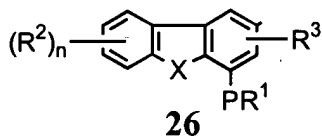


- (c) reacting the amino compound of formula (30) with ArSO_2Cl to yield a compound of formula (31)

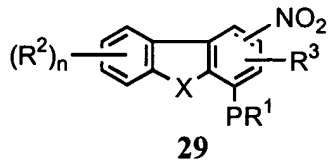


85. (New) The method of claim 62, wherein the compound of Formula (32) is prepared by:

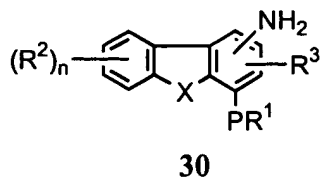
- (a) nitrating a compound of formula (26)



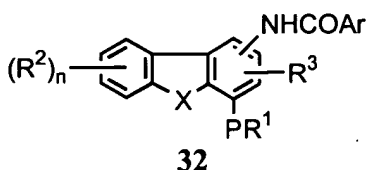
to yield a nitro compound of formula (29)



- (b) reacting the compound of formula (29) with a reducing agent to yield an amino compound of formula (30)



(c) reacting the amino compound of formula (30) with ArCOCl or a mixed anhydride of the formula ArCOOCOR^5 where R^5 is a substituted or unsubstituted alkyl, cycloalkyl, aryl, heterocyclic ring, heteroaryl, to yield a compound of formula (32)



86. (New) A compound selected from
N-(3, 5-dichloropyrid-4-yl)-4-methoxy dibenzo[b,d]furan-1-carboxamide,
N-(3, 5-dichloropyrid-4-yl)-4-methoxy dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(3,5-dichloropyrid-4-yl)-4-methoxy-8-nitro dibenzo[b,d]furan-1-carboxamide,
N-(3,5-dichloropyrid-4-yl)-4-methoxy-8-amino-dibenzo[b,d]furan-1-carboxamide, or
a pharmaceutically acceptable salt thereof.

87. (New) A compound selected from
N-(3, 5-dichloropyrid-4-yl)-4-difluoromethoxy dibenzo[b,d]furan-1-carboxamide,
N-(3,5-dichloropyrid-4-yl)-4-difluoromethoxy-dibenzo[b,d]furan-1-carboxamide-N-oxide, or
a pharmaceutically acceptable salt thereof.

88. (New) A compound selected from
N-(3,5-dichloropyrid-4-yl)-4-difluoromethoxy-8-nitro-dibenzo[b,d]furan-1-carboxamide,
N-(3,5-dichloropyrid-4-yl)-4-difluoromethoxy-8-amino-dibenzo[b,d]furan-1-carboxamide, or
a pharmaceutically acceptable salt thereof.

89. (New) A compound selected from
N-(3, 5-dichloropyrid-4-yl)-4-isopropoxy dibenzo[b,d]furan-1-carboxamide,

N-(3, 5-dichloropyrid-4-yl)-4-cyclopropylmethoxy dibenzo[b,d]furan-1-carboxamide
N-(3, 5-dichloropyrid-4-yl)-4-benzyloxy dibenzo[b,d]furan-1-carboxamide, or
a pharmaceutically acceptable salt thereof.

90. (New) A compound of claim 1 selected from

N-(pyrid-4-yl)-4-methoxy-dibenzo[b,d]furan-1-carboxamide,
N-(pyrid-4-yl)-4-methoxy-dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(2-chloropyrid-3-yl)-4-methoxy-dibenzo[b,d]furan-1-carboxamide,
N-(4-fluorophenyl)-4-methoxy-dibenzo[b,d]furan-1-carboxamide,
N-(pyrid-3-yl)-4-methoxy-dibenzo[b,d]furan-1-carboxamide,
N-(pyrid-3-yl)-4-methoxy-dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(3, 5-dichloropyrid-4-yl)-4-methoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide,
N-(3, 5-dichloropyrid-4-yl)-4-methoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(pyrid-4-yl)-4-methoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide,
N-(3, 5-dichloropyrid-4-yl)-4-difluoromethoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide,
N-(3, 5-dichloropyrid-4-yl)-4-difluoromethoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(pyrid-4-yl)-4-difluoromethoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide,
N-(pyrid-4-yl)-4-difluoromethoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(pyrid-3-yl)-4-difluoromethoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide,
N-(pyrid-3-yl)-4-difluoromethoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(pyrid-2-yl)-4-difluoromethoxy-8-trifluoromethyl dibenzo[b,d]furan-1-carboxamide,
N-(pyrid-4-yl)-4-difluoromethoxy-dibenzo[b,d]furan-1-carboxamide,
N-(pyrid-4-yl)-4-difluoromethoxy-dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(pyrid-3-yl)-4-difluoromethoxy-dibenzo[b,d]furan-1-carboxamide,
N-(pyrid-3-yl)-4-difluoromethoxy-dibenzo[b,d]furan-1-carboxamide-N1-oxide,
N-(5-chloropyrid-2-yl)-4-difluoromethoxy-dibenzo[b,d]furan-1-carboxamide, or
a pharmaceutically acceptable salt thereof.

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3,5-Dichloro-4-(4-cyclopentyloxydibenzo[b,d]-thiophen-1-ylcarboxamido)pyridine,
N1 (4-methoxyphenyl)-4-methoxydibenzo[b,d]thiophene-1-carboxamide,
N1-(4-methoxyphenyl)-4-methoxydibenzo[b,d]thiophene-1-carboxamide-5,5-dioxide,
N1-(4-chlorophenyl)-4-methoxydibenzo[b,d]thiophene-1-carboxamide,
4-(4-methoxydibenzo[b, d]thiophene-1-ylcarboxamido)pyridine,
4-(4-Cyclopentyloxydibenzo[b,d]thiophene-1-yl-carboxamido)pyridine,
3,5-Dichloro-4-(4-cyclopentyloxydibenzo[b,d]-thiophen-5,5-dioxide-1-ylcarbox-amido)pyridine-N-oxide,
3,5-Dichloro-4-(4-methoxydibenzo[b,d]-thiophen-5,5-dioxide-1-yl-carboxamido) pyridine-N-oxide,
3,5Dichloro-4-(4-methoxydibenzo[b,d]-thiophen-5,5-dioxide-1-yl-carboxamido) pyridine,
3,5 Dichloro-4-(4-difluoromethoxydibenzo[b,d]-thiophen-1-ylcarboxamido) pyridine,
N1-(4-methoxyphenyl)-4-methoxydibenzo [b,d]thiophene-1-sulfonamide,
2-(4-Methoxydibenzo[b,d] thiophen-1-ylcarboxamido)-pyridine,
4-(4-Ethoxydibenzo[b,d] thiophen-1-yl-carboxamido)-pyridine,
N1-(4-methoxyphenyl)-8,N8-dimethyl-4-methoxydibenzo[b,d]thiophen-8,1-disulfo-amide,
3-(4-Methoxydibenzo[b,d] thiophen-1-ylcarboxamido)-pyridine,
3,5-Dichloro-4-(6-ethyl-4-methoxydibenzo[b,d]-thiophen-1- ylcarboxamido)pyridine,
3,5,dichloro-4-(4-ethoxy-dibenzo[b, d]thiophen-1-yl-carboxamido)pyridine,
3-(4-Methoxydibenzo[b,d]-thiophene-5,5-dioxide-1-ylcarboxamido)-pyridine,
3,5-Dichloro-4-(4-benzyloxydibenzo[b,d]-thiophen-1- ylcarboxamido)pyridine,
N-(3,5-dichloropyrid-4-yl)-4-difluoromethoxy-8-(pyrrolidine-2-one-1-yl)-dibenzo[b,d]furan-1-carboxamide, or
a pharmaceutically acceptable salt thereof.

95. (New) A compound selected from

